

Effect of an indigenous drug formulation (churna) on clotting of mammalian blood

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Abstract

Traditional practitioners of Wedage Samaratunga Tradition of Medadumbara, Kandy, Sri Lanka uses a traditional churna consisting of ten plant ingredients (**Aralu** - *Terminalia chebula* Retz. Combretaceae, **Bulu** - *Terminalia bellerica* Roxb. Combretaceae, **Nelli** - *Emblica officinalis* Gaertn. Euphorbiaceae, **Puvak** - *Areca catechu* Linn. Palmae, **Idda** - *Wrightia zeylanica* Br. Apocynaceae, **Suduru** - *Cuminum cyminum* Linn. Umbelliferae, **Gammiris** - *Piper nigrum* Linn. Piperaceae, **Karabu** - *Eugenia caryophyllus* Thunb. Myrtaceae, **Kaippu** - *Acacia catechu* Willd. Leguminosae and **Iramusu** - *Hemidesmus indicus* R. Br. Asclepiadaceae) for the treatment of bleeding gum disorders. However, the effect of this churna on clotting of mammalian blood has not been scientifically investigated. Therefore, this study was undertaken to examine the effect of this churna on clotting of mammalian blood in vitro. Several concentrations of the extract (100%, 50%, 25%, 12.5%, 6.25%, 3.12% and 1.56%) were added to citrated goat's blood and clotting time was determined using Lee & White method. The result showed that all concentrations of the churna tested except the highest (100%) significantly shorten the clotting time ($P < 0.0001$). In contrast, the results of this study lend support for the use of this churna in low concentrations as a clotting agent in the treatment of bleeding gums.

KEYWORDS: Mammalian blood, Clotting time

Introduction

Oral hygiene is considered and established as an integral part of general health. Globally, there are about 165 million persons affected with periodontal disorders per year. Most of the periodontal disorders (gingivitis and periodontitis) are encountered in developing countries including Sri Lanka. In Sri Lanka, about 35 % of the population still rely on herbal medicine in primary health care and there are several traditional drug formulations (churnas) prescribed by Sri Lankan traditional practitioners for the treatment of gingivitis and periodontitis.

However, none of these formulations are scientifically tested for their effects on blood clotting; bleeding gums is the main clinical feature of gingivitis and periodontitis. So, we have initiated a programme to investigate the effects of locally used traditional formulations (churnas) on bleeding gum disorders on clotting of mammalian blood.

In this study, we report the effect of a churna belonging to Wedage Samaratunga tradition of Medadumbara electorate of Kandy district, Sri Lanka, on clotting of mammalian blood.

This traditional formulation (churna) consisting of ten plant ingredients (**Aralu** - *Terminalia chebula* Retz. Combretaceae, **Bulu** - *Terminalia bellerica* Roxb. Combretaceae, **Nelli** - *Emblica officinalis* Gaertn. Euphorbiaceae, **Puvak** - *Areca catechu* Linn. Palmae, **Idda** - *Wrightia zeylanica* Br. Apocynaceae, **Suduru** - *Cuminum cyminum* Linn. Umbelliferae, **Gammiris** - *Piper nigrum* Linn. Piperaceae, **Karabu** - *Eugenia caryophyllus* Thunb. Myrtaceae, **Kaippu** - *Acacia catechu* Willd. Leguminosae and **Iramusu** - *Hemidesmus indicus* R. Br. Asclepiadaceae).

Materials and methods

The plants and their parts were authenticated by Dr. M.H.A. Tissera, Wickramarachchi Ayurvedic Institute, University of Kelaniya, Sri Lanka. All ingredients except kaippu (the resin of *Acacia catechu*) are air dried on an open field for seven days. The dried materials were finely powdered (two parts of each ingredient and one part of kaippu) by using roller grinder. The testing powder stored in an airtight glass container at room temperature (30° C). 0.4g of churna was dissolved in 20ml of distilled water and thoroughly mixed. Supernatant was removed using a micropipette.

	Control	Test (Concentration of Extract)						
	0.9% Na Cl	1.56%	3.12%	6.25%	12.50%	25%	50%	100%
CT	3.52 ± 0.06	*1.78 ± 0.05	*2.10 ± 0.05	*1.87 ± 0.09	*2.15 ± 0.05	*2.15 ± 0.16	*2.00 ± 0.0	>20
CI	(3.0 - 5.0)	(1.5 - 2.0)	(2.0 - 2.5)	(1.5 - 2.0)	(2.0 - 2.5)	(1.0 - 3.0)	(2.0 - 0.0)	>20

*P < 0.0001 (Mann -Whitney U - test), n=20, CT- Clotting Time; CI - Confidence Interval

Table 1: Clotting effect of an indigenous drug formulation (Churna) on Mammalian blood
(Means ± SEM ranges in parentheses)

Citrated goat's blood was obtained from abattoir in Dematagoda, Sri Lanka. Citration was done by using 3.1% Sodium citrate. Several concentrations of the supernatant or extract (100%, 50%, 25%, 12.5%, 6.25%, 3.125%, and 1.56%) were made using appropriate amount of normal saline. For (one) 1 ml of the different concentrations of the extracts (n=20) were added to (four) 4 ml of citrated blood and mixed well.

0.2ml of 2% CaCl₂ was added to these extract - blood mixtures and the clotting times were determined at room temperature using Lee & White method.

Results are expressed as means ± SEM. Statistical analyses were made using Mann - Whitney - U test. The significance level was set at p<0.05.

Results

Results obtained are summarized in Table 1. As shown all concentrations of the extracts except 100% caused significant (p<0.0001) and marked (39% - 49%) reduction in clotting time.

Conclusions

1. The *churna* at concentrations below 50% significantly shorten the clotting time seemingly in a dose - related manner. The lowest dose is the most potent.
2. The 100% *churna* significantly prolonged the clotting time.

3. If the results are applicable to humans it justifies the use of this *churna* in low concentrations for the treatment of bleeding gum disorders.

References

1. Prevention of oral diseases, World Health Organization Publication, Geneva, 1987.
2. Wickramage, P. Conservation and sustainable development of medical plant a priority Daily news page 8.2001.
3. Bell, G, H, Davidson, N, J and Scarborough, H. Text book of Physiology and Biochemistry, E. & S. Livingston Ltd, Edinburgh and London, pp 454, 1968.
4. Ratnasooriya, W, D. and Ranatunga, K, W A plant extract that prevents clotting mamalian blood, Ceylon J. Sci (Bio-Sct.) Vol.11, No. 2, April 1975.
5. Lee, R. I and White, P. D., A clinical study of the coagulation time of Blood, Ameri. J. Med. Sci., 145,494, 1913.
6. Tocantins, L. M. and Kazal, L. A., Blood Coagulation, Haemorrhage and Thrombosis - Methods of study, Grune and Stratton, New York & London, pp. 30, 1964.
7. Biggs: Rosemary and Macfarlane, R. G. Human Blood Coagulation and its Disorders; Blackwell Scientific Publications, Oxford, pp. 380,146, 1962.